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Applicants:

Mark Ledeboer et al.

Application No.:

10/657,811

## AMENDMENTS TO THE CLAIMS

Please replace all prior versions and listings of claims with the amended claims as follows:

1. (Currently amended) A compound of formula [[I]] II or III:

or a pharmaceutically acceptable salt thereof, wherein:

R<sup>1</sup> is hydrogen or fluorine halogen,

R<sup>2</sup> is substituted or unsubstituted cycloalkyl;

each occurrence of  $R^3$  is independently <u>alkyl</u>, <u>-OH</u>, <u>-CH</u><sub>2</sub>OH or <u>alkoxy</u> <del>halogen</del>, <u>alkyl</u>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>OR<sup>4</sup>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>SR<sup>4</sup>, <u>-(CH</u><sub>1</sub>)<sub>m</sub>N(R<sup>4</sup>)<sub>2</sub>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>NR<sup>4</sup>C(O)R<sup>4</sup>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>NR<sup>4</sup>C(O)N(R<sup>4</sup>)<sub>2</sub>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>NR<sup>4</sup>CO<sub>2</sub>R<sup>4</sup>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>CO<sub>2</sub>R<sup>4</sup>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>CO<sub>2</sub>R<sup>4</sup>, <u>-(CH</u><sub>2</sub>)<sub>m</sub>SO<sub>2</sub>N(R<sup>4</sup>)<sub>2</sub>, ...

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 $\frac{(CH_2)_mS(O)R^4, \quad (CH_2)_mNR^4SO_2N(R^4)_2; \quad (CH_2)_mNR^4SO_2R^4, \quad (CH_2)_mC(=S)N(R^4)_2;}{\text{wherein m is 0, 1 or 2 and }R^4\text{ is hydrogon or alkyl};}$ 

r is 0 or [[,]] 1-or-2; and

each occurrence of R5 is independently -OH or alkyl;

p is 0 or 1; and

n is 0, 1 or 2.

- 2-4. (Canceled)
- 5. (Original) The compound of claim 1, wherein  $\mathbb{R}^1$  is F.
- 6. (Original) The compound of claim 1, wherein R<sup>1</sup> is H.
- 7. (Canceled)
- 8. (Original) The compound of claim 1, wherein n is 0.
- 9. (Original) The compound of claim 1, wherein n is 1.
- 10. (Original) The compound of c.aim 1, wherein n is 2.
- 11-14. (Canceled)
- 15. (Currently amended) The compound of claim  $\underline{1}$  3-or-4, wherein p is [[0 or]] 1 and  $R^5$  is OH, or alkyl.
- 16. (Currently amended) The compound of claim  $\underline{I}$  [[3]], wherein  $\mathbb{R}^1$  is F or H; p is 0; n is 0 or 1; r is 0 or 1; and  $\mathbb{R}^3$  is OH,  $CH_2OH$ , alkyl or alkoxy.

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17. (Currently amended) The compound of claim  $\underline{1}$  [[4]], wherein  $R^4$  is F or H; p is 0, 1 or 2; each occurrence of  $R^5$  is independently alkyl, OH,  $CH_2OH$  or alkoxy; n is 0 or 1; r is [[0 or]] 1; and  $R^3$  is OH,  $CH_2OH$ , alkyl or alkoxy.

18. (Currently amended) The compound of claim 1, wherein the compound has one of the following structures:

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- 19. (Original) A pharmaceutical composition comprising a therapeutically effective amount of a compound of claim 1 and a pharmaceutically acceptable carrier or diluent.
- 20. (Currently amended) The composition according to claim 19, further comprising an additional therapeutic agent that is selected from a treatment for stroke, a treatment for Alzheimer's Disease, a reatment for Parkinson's Disease, an agent for treating Multiple Sclerosis (MS), a treatment for asthma, an agent for treating schizophrenia, an anti-inflammatory agent, an immunomodulatory or

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immunosuppressive agent, a neurotrophic factor, an agent for treating cardiovascular disease, or an agent for treating an imm unodeficiency disorder.

- 21. (Currently amended) A method of treating a neurodegenerative, neurological, ischemic or inflammatory disorder of the central nervous system comprising administering a therapeutically effective amount of a compound of claim 1.
- 22. (Original) The method according to claim 21, wherein the ischemic disorder is stroke.
- 23. (Currently amended) The method according to claim 22, comprising the further step of:

administering to said patient an additional therapeutic agent that is selected from a treatment for stroke, a treatment for Alzheimer's Disease, a treatment for Parkinson's Disease, an agent for treating Multiple Sclerosis (MS), a treatment for asthma, an agent for treating schizophrenia, an anti-inflammatory agent, an immunomodulatory or immunosuppressive agent, a neurotrophic factor, an agent for treating cardiovascular disease, or an agent for treating an immunodeficiency disorder wherein:

said additional therapeutic agent is appropriate for the disease being treated; and

said additional therapeutic agent is administered together with said composition as a single dosage form or separately from said composition as part of a multiple dosage form.